Leishmania tarentolae contains distinct cytosolic and mitochondrial glutaminyl–tRNA synthetase activities

(aminoacyl–tRNA synthetases/trypanosomatid/mitochondrial RNA import/evolution)

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ABSTRACT The intracellular distribution of glutaminyl–tRNA synthetases and their role in mitochondrial tRNA import were evaluated in the ancient eukaryote *Leishmania tarentolae*. The following results were obtained: (*i*) Glutaminyl–tRNA synthetase was detected in leishmanial mitochondria. This was unexpected because it has been postulated that, in organelles, Gln-tRNA^{Gln} is not formed by direct acylation of tRNA^{Gln} but by enzymatic transamidation of misacylated Glu-tRNA^{Gln}. (*ii*) Whereas the cytosolic extract is able to charge cytosolic and mitochondrial tRNAs^{Gln}, the mitochondrial matrix extract does not aminoacylate the cytosol-specific tRNA^{Gln}. This indicates that mitochondrial and cytosolic glutaminyl–tRNA synthetases are distinct. (*iii*) Seven of the 11 nucleotides that differ between the cytosolic and the mitochondrial tRNA^{Gln} are sufficient to convert the cytosol-specific tRNA^{Gln} into an optimal substrate for the mitochondrial enzyme. These nucleotides are arranged in three groups consisting of the nucleotides flanking the anticodon stem, the 5′ nucleotide of the anticodon, and four nucleotides within the acceptor stem. And (*iv*), it was shown that the identity elements for recognition by the mitochondrial glutaminyl–tRNA synthetase do not overlap with a previously identified sequence segment required for mitochondrial import of the tRNA^{Gln}.

Generally, 20 functionally different aminoacyl–tRNA synthetases (aaRS), one for each amino acid, are required for protein synthesis. However, there are exceptions; in certain organisms some, tRNAs are formed by transformation of acylated tRNAs rather than by direct acylation. One example is the formation of Gln-tRNA^{Gln} because it can be achieved by two different pathways. tRNA^{Gln} can either be charged directly with glutamine by glutaminyl–tRNA synthetase (GlnRS), or formation of Gln-tRNA^{Gln} is achieved by a two-step enzymatic reaction (1). In this case, the tRNA^{Gln} is first mischarged with glutamate by glutamyl–tRNA synthetase (GluRS), resulting in Glu-tRNA^{Gln}. The glutamate attached to the tRNA^{Gln} is then converted by a specific amidotransferase into a glutamine, yielding a correctly charged tRNA^{Gln}. Absence of GlnRS and usage of the transamidation pathway to form Gln-tRNA^{Gln} has been demonstrated in a large number of organisms, including *Archea* (2), Gram-positive bacteria (3, 4), and organelles of eukaryotes (5). In agreement with this, GlnRS was lacking in *Rhizobium meliloti* (6), a member of the α-subdivision of purple bacteria and, according to the endosymbiont theory, the ancestors of present day mitochondria.

L. tarentolae belongs to the earliest diverging cells in eukaryotic evolution that have mitochondria (7). This is illustrated by a number of unique features of their mitochondria,

including a bipartite, topologically interlocked genome, guide RNA-mediated RNA editing, and the lack of mitochondrial tRNA genes (8, 9). Mitochondrial biogenesis in trypanosomatids therefore not only involves import of proteins, as in all other eukaryotes, but also import of the whole set of nuclearly encoded mitochondrial tRNAs (10–13). However, mitochondrial tRNA import is not restricted to protozoa but has also been shown in plants and yeast (14, 15). aaRS have been implicated in the import process. Whereas there is only indirect evidence for their involvement in plants (16), participation of a precursor of a mitochondrial aaRS in tRNA import has been shown directly in yeast (17). In *L. tarentolae*, in most cases, one gene codes for both mitochondrial and cytosolic tRNA isotypes, resulting in almost identical sets of tRNAs in the two compartments (10). This raises the questions whether, in *L. tarentolae*, the aaRS as well are identical in the cytosol and the mitochondrial fraction and whether these enzymes play a role in tRNA import.

To address these questions, we focused on the tRNAs^{Gln}. Two highly homologous tRNAs^{Gln} have been characterized in *L. tarentolae*: the tRNA^{Gln} with the anticodon UUG, which, like most other leishmanial tRNAs is found in the cytosol as well as in mitochondria (18), and the tRNA^{Gln} with the anticodon CUG, which is the only known cytosol-specific tRNA in that organism (19). The intracellular distribution and the substrate specificities of GlnRSs were determined. Unexpectedly, GlnRS activity was found in mitochondrial matrix fractions of *L. tarentolae*, showing that GlnRS is not universally absent from mitochondria. In addition, it was shown that the mitochondrial GlnRS of *L. tarentolae* is distinct from its cytosolic counterpart. Furthermore, it could be demonstrated that the sequence elements within mitochondrial tRNA^{Gln} that are identity determinants for the mitochondrial GlnRS are distinct from a previously identified import signal in the D loop (20).

MATERIALS AND METHODS

Cells. *L. tarentolae* (UC strain) was grown at 27°C in Difco brain heart infusion medium containing 10 mg/ml hemin to late log phase (0.5×10^8–1.5×10^8 cells/ml) and used immediately. Cells were washed once in cold 20 mM sodium-phosphate buffer (pH 7.9) containing 150 mM NaCl and 20 mM glucose.

Isolation of tRNA. Washed *L. tarentolae* (3×10^{11} cells) were resuspended in 10 ml of 20 mM Tris·HCl/20 mM magnesium acetate, pH 7.4. Cells were extracted twice with an equal volume of phenol and equilibrated in 25 mM sodium acetate (pH 4.5) containing 50 mM NaCl. Sodium acetate (pH 5.2) was added to 0.3 M and total RNAs were precipitated by the addition of an equal volume of isopropanol. The RNA pellet

Abbreviations: aaRS, aminoacyl–tRNA synthetase; GlnRS, glutaminyl–tRNA synthetase; GluRS, glutamyl–tRNA synthetase.
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was resuspended in 4 ml of 0.2 M Tris·acetate (pH 9.0) and deacylated for 30 min at 37°C. After addition of sodium acetate to 0.3 M, the RNAs were precipitated by the addition of 2 vol of ethanol. The resulting pellet was resuspended in 2 ml of 10 mM 1,3-bis[tris(hydroxymethyl)methylamino]propane·HCl (pH 7.0), 1 mM MgCl$_2$, and 1 mM 1,4-dithiothreitol and treated with 10 units of RQ1 DNase (Promega) for 20 min at 37°C. Subsequently, the mixture was phenol-extracted and precipitated with ethanol as before. The pellet was dissolved in 3 ml of 1 M NaCl and centrifuged for 10 min at $16,000 \times g$ to remove high molecular weight RNAs. The supernatant containing the tRNAs was ethanol precipitated, resuspended in 1 ml of H$_2$O, and loaded on a 0.6 ml DEAE-cellulose column equilibrated in buffer A (10 mM sodium acetate/0.2 M NaCl/10 mM MgCl$_2$, pH 5). The column was washed with 4 ml of buffer A and eluted by 2 ml of buffer A containing 0.7 M NaCl. The tRNAs (yield 2–3 mg) were precipitated, resuspended in H$_2$O, and frozen at -70°C until further use.

***In Vitro* Transcription of tRNA.** The tRNA$^{\text{Gln}}$ variants were produced by *in vitro* transcription using T7 polymerase (21). DNA-fragments containing the T7 promoter and the corresponding natural or synthetic tRNA$^{\text{Gln}}$ genes were produced by PCR using the following oligonucleotides: QM1, 5′-GGAATTCT<u>AATACGACTCACTATAGGTC</u>-CTATAGTGTAGT-3′; QM2, 5′-TGGTGGTCCTACCAG-GAT-3′; QC1, 5′-GGAATTC<u>TAATACGACTCACTAT-AGCTCCTATAGTGTAGCGG</u>-3′; QS1, 5′-<u>TAATACGA-CTCACTATAG</u>CTCCTATAGTCTAGTTGGTTAGGA-CCTCGG ACTCTGAAT C-3′; QCUUG, 5′-<u>TAATAC-GACTCACTATAG</u>CTCCTATAGTGTAGCG-GTTATCACCTCGGACTTTGA ATCCGAT-3′; QCM1, 5′-<u>TAATACGACTCACTATAGGTCCTATAGT</u>-GTAGCGGTT-3′; ACST, 5′-<u>TAATACGACTCACTATAG</u>GTCCTATAGTGTAGTGGTTATCACT-3′; QC2, 5′-TG-GCACTCCTACCTGGACTC-3′; QCM2, 5′-TGGTGGTC-CTACCTGGACTCGAAC-3′; ACST-3′, 5′-TGGTGGTC-CTACCTGGACTCGAACCAGGGTTT-3′; GNLT1, 5′-GCGGTACCGATGTTATGGGCTAACCG-3′; GNLT2, 5′-GGAGATCTCCCCAGCCGCTTTTTGG-3′; LTQM1, 5′-CTAGATCACCCACGCTTG-3′; and LTQM2, 5′-GAG-TGCCGTACTCCCACT-3′. Sequences corresponding to the T7 promoter are underlined. As DNA templates to produce the variant tRNAs$^{\text{Gln}}$, in addition to some of the PCR products listed below, the following PCR products were used: PCR-fragment GlnCyt was produced using the primers GNLT1/ GNLT2 and genomic leishmanial DNA as template; PCR-fragment GlnMit was produced using the primers LTQM1/ LTQM2 and genomic leishmanial DNA as template. The variant tRNA$^{\text{Gln}}$ genes containing a 5′-flanking T7 promoter were prepared using the following primer pairs and templates: cytosolic tRNA$^{\text{Gln}}$ (QC1/QC2, template GlnCyt); cytosolic tRNA$^{\text{Gln}}$ containing the mitochondrial acceptor stem (QCM1/QCM2, template PCR-product of the previous reaction); cytosolic tRNA$^{\text{Gln}}$ containing the mitochondrial anticodon (QCUUG/QC2, template GlnCyt); cytosolic tRNA$^{\text{Gln}}$ containing the mitochondrial acceptor stem and the anticodon (QCM1/QCM2, template PCR product of the previous reaction); cytosolic tRNA$^{\text{Gln}}$ containing the mitochondrial acceptor stem and anticodon, the nucleotide substitutions flanking the anticodon stem and the D loop substitution (ACST/ ACST3′, template GlnMit); cytosolic tRNA$^{\text{Gln}}$ containing all the previous changes except the D loop substitutions (QCM1/ ACST3′, template PCR product of previous reaction); mitochondrial tRNA$^{\text{Gln}}$ (QM1/QM2, template GlnMit); and tRNA$^{\text{Gln}}$(D-Ile) (QS1/QC2, template GlnCyt).

Preparation of RNA-Free Cytosolic Fractions. Washed *L. tarentolae* cells were hypotonically lysed at 2.4×10^9 cells/ml in 1 mM Tris·HCl/1 mM EDTA, pH 8, by four passages through a 25-G syringe needle. A quarter volume of 5X acylation buffer (250 mM Tris·HCl, pH 7.5/125 mM KCl/40

mM MgCl$_2$/50 mM 2-mercaptoethanol), one-ninth volume of 100% glycerol was added, and the extract was centrifuged for 25 min at $150,000 \times g$ at 4°C. NaCl was added to 150 mM, and the sample was applied on a DEAE cellulose column (13 mg protein/ml bed volume) equilibrated in the same buffer. The flow through fraction was concentrated using a Centricon-30 concentrator (Amicon) to 1 ml and loaded on a 10 ml Sephadex G-100 column (20 cm length) equilibrated in 1x acylation buffer containing 10% glycerol (22). Protein-containing fractions of the exclusion volume were pooled (yield 2.3 mg/10^{10} cells), frozen in liquid N$_2$, and stored at -70°C until further use in the charging assays.

Preparation of RNA-Free Mitochondrial Matrix Fractions. Mitoplasts of *L. tarentolae* were purified as described (23, 24). Isolated mitoplasts (40 mg protein) were resuspended in 20 mM Tris·HCl (pH 8), 250 mM sucrose, and 2 mM EDTA and frozen in liquid N$_2$. To prepare the mitochondrial matrix fraction, an equal volume of 16 mM MgCl$_2$, 100 mM Tris·HCl (pH 7.5), and 50 mM KCl containing 2 mM phenylmethylsulfonylfluoride was added to thawed mitoplasts, and the suspension was sonicated on wet ice (10 bursts of 5 s using the microtip of a sonicator (MSE, Zivy, Switzerland), amplitude 3 at medium power setting). The extract was then subjected to another freeze thaw cycle, and the sonication step was repeated. After the addition of glycerol to 10% and 2-mercaptoethanol to 10 mM, the suspension was centrifuged at $180,000 \times g$ at 4°C for 25 min. Approximately 35% of total proteins were recovered in the supernatant corresponding to the mitochondrial matrix fraction. The supernatant was adjusted to 150 mM NaCl and applied to a DEAE cellulose (12 mg matrix protein/0.5 ml bed volume) followed by Sephadex G-100 chromatography as described for the cytosol (see previous paragraph). Approximately 2.5 mg of mitochondrial matrix proteins was recovered in the final fraction.

Yeast mitochondria originating from the wild-type strain D273–10B were purified using Nycodenz gradients (25). Mitochondrial RNAs were isolated as described (26). From 20 mg of mitochondria, 270 μg of RNA was obtained. RNA-free mitochondrial matrix fraction was prepared as described for *L. tarentolae*. Fifty milligrams of purified mitochondria yielded 4 mg of RNA-free matrix fraction.

Charging Assays. Aminoacylation reactions for *L. tarentolae* were performed in 100 μl of 1x acylation buffer supplemented with 4 mM ATP containing 20 μg of total leishmanial tRNA or 1.8 μg of the corresponding *in vitro* transcribed tRNA, 60 μg of crude cytosolic or mitochondrial aaRS fraction and 19 μM of ^{14}C-glutamine (258 mCi/mmol) or ^{14}C-glutamate (265 mCi/mmol) (Dupont) respectively. For the competition assays an additional 1.8 or 9 μg of the competitor tRNA [cytosolic tRNA$^{\text{Gln}}$ or tRNA$^{\text{Gln}}$(D-Ile)] were added to a standard charging reaction. Charging assays for yeast were done under identical conditions with the exception that 50 μg of total mitochondrial RNA and 100 μg of crude mitochondrial aaRS-containing matrix fraction were used. All samples were incubated for 15 min at 37°C and spotted onto dried Whatman filter paper (2.5 x 2.5 cm) which had been pretreated with 5% TCA containing either 0.1 mM glutamine or glutamate. The filters were washed three times (15 min each) on ice in 4 ml/filter of cold 5% TCA containing 0.1 mM glutamine or glutamate followed by two washes in cold 0.1 M HCl. After a final wash in cold ethanol (96%) the filters were dried and processed for scintillation counting.

Thin Layer Chromatography. After incubation at 37°C, a standard charging reaction containing total tRNAs, leishmanial mitochondrial matrix extract, and ^{14}C-glutamine was phenol-extracted, ethanol-precipitated, and washed with 75% ethanol. The resulting pellet was deacylated in 50 μl of 10 mM KOH for 10 min at 65°C. Subsequently, the pH was adjusted (pH 6–7) by the addition of 0.1 M HCl and dried under vacuum. Finally, the pellet was resuspended in 3 μl of water

and spotted onto a cellulose thin layer chromatography plate. The plate was developed for ≈3 h in a mixture of isopropanol/ formic acid/water (77:18.2:4.8) and exposed on x-ray film.

Miscellaneous. Antiserum directed against yeast Hsp60 was a generous gift of G. Schatz, Biozentrum, Basel. Polyclonal antiserum specific for leishmanial pyruvate kinase was kindly provided by P. Michels, International Institute of Cellular and Molecular Pathology, Brussels. Both sera were used at a dilution of 1/500. The signals were visualized using peroxi-dase-conjugated second antibody and the enhanced chemilu-minescence detection kit (Amersham). Protein concentrations were determined using the BCA assay (Pierce).

RESULTS

Mitochondrial GlnRS in *L. tarentolae*. *In vitro* charging assays using labeled glutamine or glutamate were performed to measure GlnRS and GluRS activities in the mitochondria and the cytosol of *L. tarentolae*. Because of the nuclear origin of mitochondrial tRNAs in *L. tarentolae*, their cytosolic and mitochondrial sets of tRNAs overlap to a great extent (10). Therefore, tRNAs isolated from total cellular extract were used as substrates for both cytosolic and mitochondrial aaRS. Similar specific activities of GlnRS and GluRS were detected in the cytosol (Fig. 1*A*, left). Unexpectedly, not only GluRS but also GlnRS activity was detected in the corresponding mito-chondrial matrix fractions (Fig. 1*A*, right). In yeast mitochon-dria assayed under the same conditions, however, only GluRS activity could be measured (Fig. 1*B*). Cytosolic contamination of the mitochondrial fraction, a possible trivial explanation for our results, can be excluded for the following reasons: (*i*) Immunoblots using antibodies against the mitochondrial heat shock protein 60 and the cytosolic protein pyruvate kinase demonstrate that very little cytosolic contamination is ob-served in the mitochondrial fraction (Fig. 1*C*). (*ii*) Comparable specific activities of cytosolic and mitochondrial GlnRS are detected. Should GlnRS be a cytosolic contaminant of the mitochondrial fraction, we would expect much lower specific activity. (*iii*) Comparable specific activities are observed for GlnRS and GluRS (Fig. 1*A*), indicating that, in *L. tarentolae*, GlnRS behaves like a bona fide mitochondrial aaRS. In the aminoacylation assays, crude mitochondrial matrix fraction was used so that metabolic conversion of glutamine to gluta-mate before charging could not be excluded. In this case, mitochondrial GluRS but not GlnRS activity would have been measured in the assay, even though labeled glutamine and not glutamate was added to the reaction. To rule out this possi-bility, we performed a preparative *in vitro* charging experiment using radioactive glutamine, total tRNA, and mitochondrial matrix fraction as a source of GlnRS activity. After the reaction, the charged tRNAs were precipitated and deacy-lated. The labeled amino acid released in this process was analyzed by thin layer chromatography and shown to comi-grate with glutamine but not with glutamate (Fig. 1*D*), dem-onstrating that bona fide GlnRS activity was measured in the assay.

In vitro charging experiments also were performed using the mitochondrial matrix fraction of *Trypanosoma brucei*. Qualitatively identical results as for *L. tarentolae* were obtained. Similar levels of activity of GlnRS and GluRS were measured in the mitochondrial matrix of trypanosomes (data not shown); however, the specific activities were significantly lower than in *Leishmania*. These results demonstrate the existence of GlnRS in the mitochondria of trypanosomatids, indicating that this enzyme is not universally absent in mitochondria.

Cytosolic and Mitochondrial GlnRS Are Distinct. Two tRNAs[Gln] have been characterized in *L. tarentolae*. They differ in 11 nucleotide positions and in their intracellular localization (Fig. 2*A*). The tRNA[Gln](CUG), unlike most other leishmanial
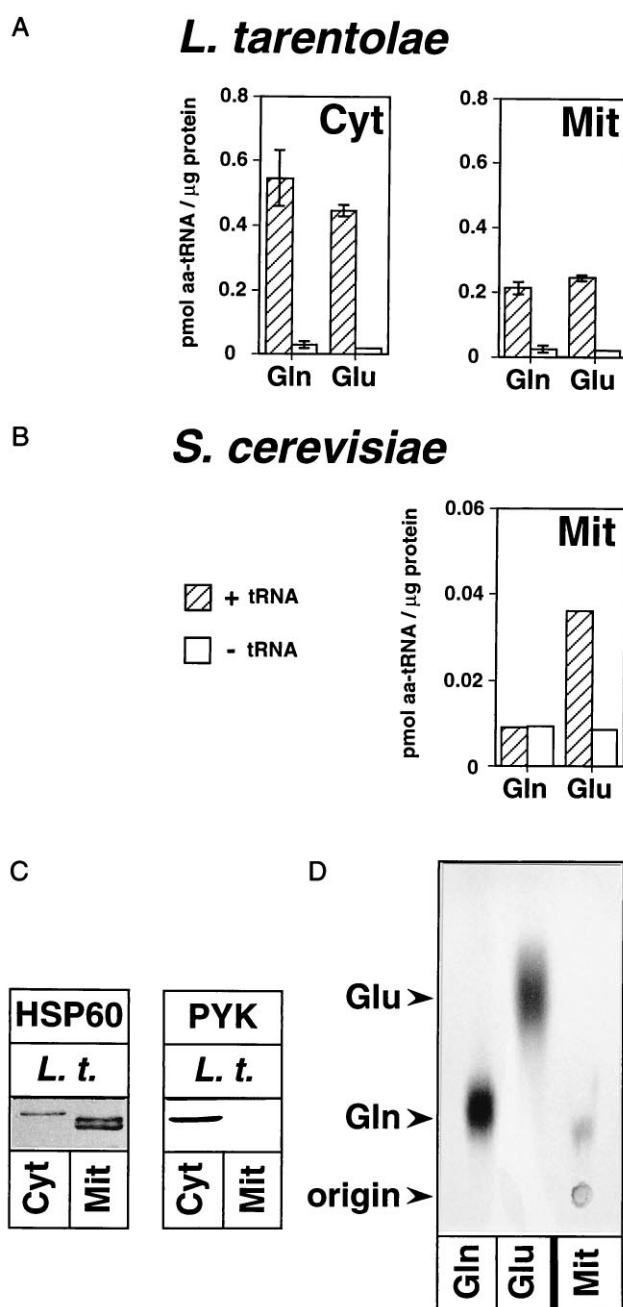
FIG. 1. Mitochondrial GlnRS in *L. tarentolae*. (*A*) Leishmanial total tRNAs were charged with ^{14}C-glutamine (Gln) or ^{14}C-glutamate (Glu) using 60 μg each of RNA-depleted cytosolic (Cyt) or mitochondrial matrix (Mit) fractions of *L. tarentolae*. Bars indicate mean values (±SD) of independent aminoacylation reac-tions (*n* = 4–8) using at least two independently prepared cytosolic or mitochondrial extracts. (*B*) Yeast mitochondrial tRNA was charged with ^{14}C-glutamine (Gln) or ^{14}C-glutamate (Glu) using 100 μg of RNA-depleted yeast mitochondrial matrix extract. The reac-tions were incubated in the presence (hatched columns) or as a control in the absence of the corresponding tRNAs (empty col-umns). The *y* axis indicates the specific activities corresponding to picomolars of aminoacylated tRNA formed per μg of proteins. (*C*) Equal amounts (50 μg) of RNA-depleted cytosolic (Cyt) and mitochondrial matrix (Mit) fraction used for the charging experi-ments were analyzed by immunoblots using antisera specific for mitochondrial heat shock protein 60 (HSP60) or the cytosolic protein pyruvate kinase (PYK). (*D*) A charging reaction containing total tRNAs, ^{14}C-glutamine, and mitochondrial matrix extract was deacylated, and the released amino acid (Mit) was analyzed by thin layer chromatography. ^{14}C-glutamine (Gln) and ^{14}C-glutamate (Glu) served as markers.
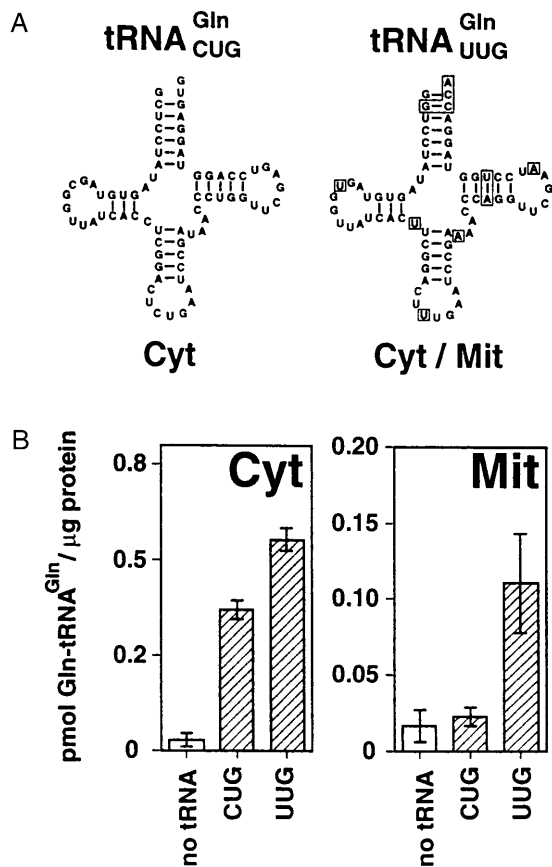
FIG. 2. Cytosolic and mitochondrial GlnRS are distinct. (*A*) Inferred secondary strucures of cytosol-specific (Cyt) and mitochondrial (Cyt/Mit) tRNAs[Gln] of *L. tarentolae*. Nucleotides in the mitochondrial tRNA[Gln] that are different from the cytosolic ones are boxed. (*B*) Cytosol-specific tRNA[Gln] (CUG) and mitochondrial tRNA[Gln] (UUG) of *L. tarentolae* were charged with ^{14}C-glutamine using 60 μg each of RNA-depleted cytosolic (Cyt) or mitochondrial matrix (Mit) fractions. As a control, the reactions were performed without adding the tRNAs (no tRNA). Bars indicate mean values (±SD) of independent aminoacylation reactions (*n* = 4–8) using at least two independently prepared cytosolic or mitochondrial extracts. *Y* axis as Fig. 1.

tRNAs, is found exclusively in the cytosol (19). The tRNA[Gln] (UUG), however, has a dual location and is found in both the cytosol and the mitochondria (18). We will refer in this study to the tRNA[Gln](CUG) as the cytosolic and the tRNA[Gln]-(UUG) as the imported mitochondrial tRNA[Gln], even though the latter also is found in the cytosol. *In vitro* charging assays were performed using *in vitro*-transcribed cytosolic or mitochondrial tRNAs[Gln] as substrates and cytosolic or mitochondrial extract as source of enzyme. Cytosolic extract aminoacylated both substrates whereas the mitochondrial extract was only able to charge the mitochondrial but not the cytosolic tRNA[Gln] (Fig. 2*B*). These results prove that the mitochondrial GlnRS activity cannot be caused by cytosolic contamination of the mitochondrial fraction, and, most importantly, they show that cytosolic and mitochondrial GlnRS activities are distinct.

Identity Elements Recognized by Mitochondrial GlnRS. Cytosolic and mitochondrial tRNA[Gln] differ by 11 nucleotides that are distributed in six groups all over the molecule (Fig. 2*A*). To determine which groups of nucleotides, or which combination of groups, are critical for recognition by the mitochondrial GlnRS, the cytosolic tRNA[Gln] sequences were successively replaced by the corresponding mitochondrial tRNA[Gln] sequences (Fig. 3). The resulting hybrid molecules corresponded to the cytosolic tRNA[Gln] containing (*i*) the mitochondrial acceptor stem, (*ii*) the mitochondrial anticodon,
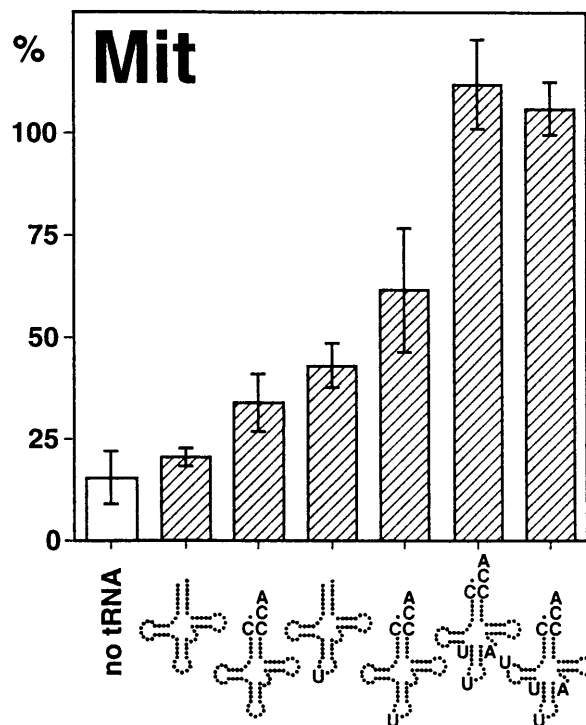

FIG. 3. Identity elements recognized by mitochondrial GlnRS. Wild-type cytosolic tRNA[Gln] and cytosolic tRNA[Gln] variants were charged with ^{14}C-glutamine using 60 μg of RNA-depleted mitochondrial matrix (Mit) fractions. The tested tRNA[Gln] variants are indicated along the *x* axis. Nucleotides in the cytosol-specific tRNA[Gln] that have been changed to the mitochondrial ones are indicated by letters. The level of activity of each tRNA[Gln] variant is expressed relative to the charging levels reached by mitochondrial tRNA[Gln]. Bars indicate mean values (±SD) of independent aminoacylation reactions (*n* = 3–8) using two independently prepared mitochondrial extracts. The background value of a charging reaction without added tRNA (no tRNA) also is shown.

(*iii*) the mitochondrial acceptor stem and the anticodon, (*iv*) the mitochondrial acceptor stem, the anticodon, and the nucleotide substitutions flanking the anticodon stem, and (*v*) all of the previous changes and the D loop substitution. *In vitro* transcripts of all of these tRNA[Gln] variants were tested for charging by the mitochondrial extract. Three elements required for recognition of the mitochondrial GlnRS were identified: (1) the nucleotides flanking the anticodon stem that contribute approximately half and (2) the 5′ nucleotide of the anticodon and (3) the acceptor stem each contributing approximately one-fourth to the conversion of the cytosolic tRNA[Gln] into a substrate for the mitochondrial GlnRS. If all three elements are present, the same level of activity was reached as with the natural substrate, the mitochondrial tRNA[Gln]. The nucleotide substitutions in the D loop or the T loop therefore are not identity elements.

Import Signal and Identity Elements for Mitochondrial GlnRS Are Distinct. Mitochondrial tRNA[Gln] must not only contain recognition elements for the mitochondrial GlnRS but also elements that specify its import into mitochondria. Recently Lima *et al.* (20) showed that, in *L. tarentolae*, the cytosolic tRNA[Gln] is converted into an imported tRNA if its D stem loop is replaced with that of tRNA[Ile], which is predominantly found in mitochondria. The five point mutations necessary to switch the tRNA[Gln] D stem loop to the tRNA[Ile] D stem loop are indicated in Fig. 4*A*; the resulting molecule was called tRNA[Gln](D-Ile). From these results, it was concluded that at least one import signal must lie within the D loop region.
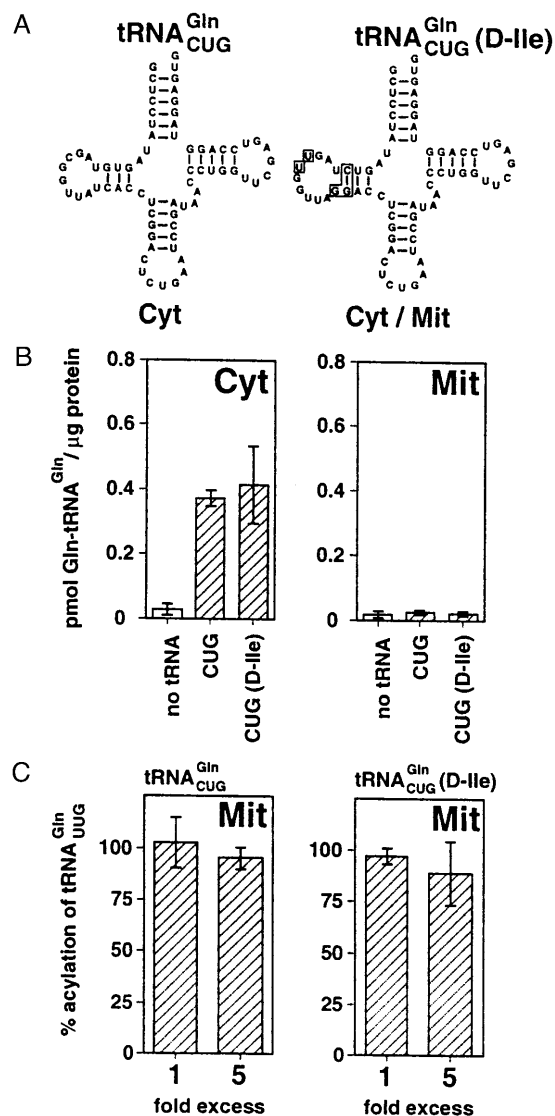
FIG. 4. Import signal and identity elements for mitochondrial GlnRS are distinct. (*A*) Inferred secondary strucures of cytosol-specific tRNA[Gln] (Cyt) and mutant tRNA[Gln], which is imported into mitochondria (Cyt/Mit) (20). Nucleotides in the tRNA[Gln](D-Ile) that are different from the cytosolic ones are boxed. (*B*) Cytosol-specific tRNA[Gln] (CUG) and mutant imported tRNA[Gln] [CUG(D-Ile)] were charged with ^{14}C-glutamine using 60 μg each of RNA-depleted cytosolic (Cyt) or mitochondrial matrix (Mit) fractions. As a control, the reactions were performed without adding tRNAs (no tRNA). *Y* axis as Fig. 1. (*C*) Competition assays were performed by charging of mitochondrial tRNA[Gln] (UUG) with ^{14}C-glutamine using 60 μg of mitochondrial matrix (Mit) in the presence of a 1- or 5-fold excess of cytosolic tRNA[Gln] (CUG) (*Left*) or the mutant-imported tRNA[Gln] [CUG(D-Ile)] (*Right*). The level of charging is expressed as percentage of a standard charging reaction without competitor tRNA. Bars indicate mean values (±SD) of independent aminoacylation reactions (*n* = 3–8) using at least two independently prepared cytosolic or mitochondrial extracts.

All mitochondrial aaRS are imported into mitochondria. It has therefore been suggested that these enzymes may mediate mitochondrial tRNA import (27). Indeed, direct involvement of an aaRS in mitochondrial tRNA import has been shown in yeast (17). In addition, it was shown in plants that a point mutation within the tRNA[Ala] that abolishes charging also prevents its import into mitochondria, providing evidence that aaRS are also involved in tRNA import in plants (16). To assess the role mitochondrial GlnRS might play in tRNA import in *L. tarentolae*, the tRNA[Gln](D-Ile) was transcribed *in vitro* and

subjected to charging assays. The tRNA[Gln](D-Ile) behaved identically to the cytosolic tRNA[Gln], in that both could be charged by the cytosolic but not by the mitochondrial GlnRS (Fig. 4*B*). Cytosolic tRNA[Gln] and tRNA[Gln](D-Ile), however, differ in their intracellular localization; only tRNA[Gln](D-Ile) is imported into mitochondria, indicating that charging by mito-chondrial GlnRS is not required for import of tRNA[Gln](D-Ile). Nevertheless, the fact that charging and mitochondrial import tRNA[Gln](D-Ile) are not coupled does not preclude the in-volvement of mitochondrial GlnRS in import because the imported tRNA[Gln](D-Ile) may still interact with the enzyme without being charged. Because the mitochondrial GlnRS has not been purified, direct binding of tRNA[Gln](D-Ile) to GlnRS cannot be assayed. However, should such a binding occur, it is possible to test whether it involves the same domain of GlnRS that is required for aminoacylation. In this case, tRNA[Gln](D-Ile), which is imported but not charged, is expected to interfere with aminoacylation of mitochondrial tRNA[Gln] by mitochon-drial GlnRS. Fig. 4*C* demonstrates that neither cytosolic tRNA[Gln], which is not expected to bind to the enzyme and therefore serves as a control, nor tRNA[Gln](D-Ile) inhibited charging of mitochondrial tRNA[Gln] by GlnRS when added to the reaction in up to a 5-fold excess. It is therefore concluded that no significant binding of tRNA[Gln] to the domain of mitochondrial GlnRS, which is responsible for charging, is required for import. Mitochondrial GlnRS might, however, still be involved in import. In this case, though, one must postulate that it contains two distinct nonoverlapping binding sites for imported tRNA[Gln], one for charging and one for import.

DISCUSSION

There is overwhelming evidence that mitochondria arose from endosymbiontic capture of a member of the α-subdivision of purple bacteria. It is generally assumed that the endosymbi-ontic event happened just once in eukaryotic evolution and that therefore all mitochondria are evolutionarily related (28). Recently, it was shown that *R. meliloti* (6), a member of the α-subdivision of purple bacteria, unlike other Gram-negative bacteria, uses the transamidation pathway for Gln-tRNA[Gln] formation (6). In line with this, it was demonstrated also that mitochondria use the transamidation pathway like their pre-sumed ancestors and therefore are devoid of mitochondrial GlnRS (5). The transamidation pathway is therefore consid-ered to be a more ancestral pathway to form Gln-tRNA[Gln] than direct acylation by GlnRS.

Using *in vitro* charging assays with radioactive glutamine and RNA-free cytosolic and mitochondrial extracts, we show here that GlnRS is not universally absent from mitochondria but is found in organelles of *L. tarentolae* and *T. brucei*. Evidence is presented that shows that, in *L. tarentolae*, mitochondrial and cytosolic GlnRS are distinct because they exhibit a different substrate specificity. One interpretation of these results is that the cytosolic GlnRS is able to charge both cytosolic and mitochondrial tRNAs[Gln]. The mitochondrial GlnRS, which has a more limited substrate specificity, would in this scenario be localized in mitochondria only. However, it cannot be excluded that cytosolic and mitochondrial GlnRS have nonoverlapping substrate specificities. In this case, one needs to postulate a dual location of the mitochondrial enzyme, in both cytosol and mitochondria.

GlnRS is not found in the α-subdivision of purple bacteria, the presumed ancestors of present day mitochondria (6). Could this mean that trypanosomatid mitochondria are derived from a different endosymbiont than other mitochondria? Indeed, mitochondria of trypanosomatids are unique in other respects as well. Their genome consists of a complex intercalated network of two genetic elements, the maxi- and the minicircles (29). Nowhere else in nature has a comparable structure been

found. In addition, many mitochondrial genes in trypanosomatids represent cryptogenes, whose transcripts have to be extensively edited to become functional mRNAs (9). RNA editing was shown to be mediated by short RNAs, called guide RNAs (30), which have not been found in mitochondria of any other organism. These features would argue for a polyphyletic origin of mitochondria. However, a monophyletic origin of mitochondria is favored by the fact that the mitochondrial genes in trypanosomes code for the same set of proteins as most other mitochondria. In this case, mitochondrial GlnRS would have been acquired later in evolution after the endosymbiontic event. This could have been achieved by gene duplication of the cytosolic GlnRS and establishing of a mitochondrial targeting sequence. At present, it is not possible to decide whether mitochondrial GlnRS in trypanosomatids has been acquired early or late in evolution. Cloning and sequencing of mitochondrial GlnRS might help to answer this question.

It is interesting to note that mitochondrial GlnRS may also exist in other protozoa. Already more than 20 years ago, GlnRS activity was measured in *Tetrahymena* mitochondria (27). This work was done before the discovery of transamidation as the primary pathway to form Gln-tRNA^{Gln} in mitochondria, however, and therefore needs to be confirmed. The reason why trypanosomatid and maybe other protozoal mitochondria contain GlnRS and do not use the transamidation pathway is unknown at present. Of interest, both trypanosomatids and *Tetrahymena* (31, 32) import all or most of their mitochondrial tRNAs, including the tRNAs^{Gln}, from the cytosol. However, we were able to show that, in *L. tarentolae*, there is no direct link between charging by mitochondrial GlnRS and import of tRNA^{Gln}. This conclusion is based on the identification of the sequence elements necessary for charging by mitochondrial GlnRS, which can be separated from a previously identified import signal located in the D loop of the tRNA (20). Consequently, a mutated tRNA^{Gln} that cannot be charged by mitochondrial GlnRS can still be imported into mitochondria, provided that it contains an import signal in the D loop. This is in contrast with the plant system, in which a single point mutation within the acceptor stem concomitantly inactivated both charging by the cognate aaRS and mitochondrial import (16). In yeast, the situation is different. Mitochondrial import of the single imported tRNA^{Lys} is mediated by the mitochondrial precursor of lysyl–tRNA synthetase. Surprisingly, this enzyme is only able to bind, but not to aminoacylate, the imported tRNA^{Lys} (17). The same could potentially be true in *L. tarentolae*. However, neither an excess of cytosolic tRNA^{Gln} nor an excess of import competent tRNA^{Gln}(D-Ile) was able to inhibit charging of mitochondrial tRNA^{Gln} by mitochondrial extracts, indicating that the binding sites for charging and the putative binding sites for import do not overlap. In addition, it has been shown in *T. brucei* that intron-containing tRNA^{Tyr} that could not be charged could still be imported into mitochondria *in vivo* (33). These results, together with the observation that *in vitro* import of tRNA into mitochondria of *Leishmania donovani* does not require cytosolic factors (34, 35), make the involvement of aaRS in mitochondrial tRNA import in trypanosomatids unlikely.

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